

**BICALUTAMIDE- bicalutamide tablet, film coated**  
**Mylan Pharmaceuticals Inc**

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**HIGHLIGHTS OF PRESCRIBING INFORMATION**

**These highlights do not include all the information needed to use bicalutamide tablets safely and effectively. See full prescribing information for bicalutamide tablets.**

**BICALUTAMIDE tablets, USP for oral use**

**Initial U.S. Approval: 1995**

----- **RECENT MAJOR CHANGES** -----

Dosage and Administration (2.1) 02/2015

----- **INDICATIONS AND USAGE** -----

- Bicalutamide tablets, USP 50 mg are an androgen receptor inhibitor indicated for use in combination therapy with a luteinizing hormone-releasing hormone (LHRH) analog for the treatment of Stage D<sub>2</sub> metastatic carcinoma of the prostate.
- Bicalutamide tablets, USP 150 mg daily are not approved for use alone or with other treatments. (1)

----- **DOSAGE AND ADMINISTRATION** -----

The recommended dose for bicalutamide tablets therapy in combination with an LHRH analog is one 50 mg tablet once daily (morning or evening). (2)

----- **DOSAGE FORMS AND STRENGTHS** -----

50 mg tablets (3)

----- **CONTRAINDICATIONS** -----

- Hypersensitivity (4.1)
- Women (4.2)
- Pregnancy (4.3 and 8.1)

----- **WARNINGS AND PRECAUTIONS** -----

- Severe hepatic injury and fatal hepatic failure have been observed. Monitor serum transaminase levels prior to starting treatment with bicalutamide tablets, at regular intervals for the first 4 months of treatment and periodically thereafter, and for symptoms or signs suggestive of hepatic dysfunction. Use bicalutamide tablets with caution in patients with hepatic impairment. (5.1)
- Gynecomastia and breast pain have been reported during treatment with bicalutamide tablets 150 mg when used as a single agent. (5.2)
- Bicalutamide tablets are used in combination with a LHRH agonist. LHRH agonists have been shown to cause a reduction in glucose tolerance in males. Consideration should be given to monitoring blood glucose in patients receiving bicalutamide tablets in combination with LHRH agonists. (5.3)
- Monitoring Prostate Specific Antigen (PSA) is recommended. Evaluate for clinical progression if PSA increases. (5.4)

----- **ADVERSE REACTIONS** -----

Adverse reactions that occurred in more than 10% of patients receiving bicalutamide tablets plus an LHRH-A were: hot flashes, pain (including general, back, pelvic and abdominal), asthenia, constipation, infection, nausea, peripheral edema, dyspnea, diarrhea, hematuria, nocturia and anemia. (6.1)

**To report SUSPECTED ADVERSE REACTIONS, contact Mylan Pharmaceuticals Inc. at 1-877-446-3679 (1-877-4-INFO-RX) or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch).**

----- **DRUG INTERACTIONS** -----

- R-bicalutamide is an inhibitor of CYP 3A4; therefore, caution should be used when bicalutamide tablets are co-administered with CYP 3A4 substrates. (7)
- Prothrombin times should be closely monitored in patients already receiving coumarin anticoagulants who are started on bicalutamide tablets. (7)

----- **USE IN SPECIFIC POPULATIONS** -----

- Pediatric patients: Efficacy has not been demonstrated for the treatment of familial male-limited precocious puberty

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## FULL PRESCRIBING INFORMATION

### 1 INDICATIONS AND USAGE

**Bicalutamide tablets USP, 50 mg daily are indicated** for use in combination therapy with a luteinizing hormone-releasing hormone (LHRH) analog for the treatment of Stage D<sub>2</sub> metastatic carcinoma of the prostate.

**Bicalutamide tablets USP, 150 mg daily are not approved** for use alone or with other treatments [*see Clinical Studies (14.2)*].

### 2 DOSAGE AND ADMINISTRATION

#### 2.1 Recommended Dose and Schedule

The recommended dose for bicalutamide tablets therapy in combination with an LHRH analog is one 50 mg tablet once daily (morning or evening), with or without food. It is recommended that bicalutamide tablets be taken at the same time each day. Treatment with bicalutamide tablets should be started at the same time as treatment with an LHRH analog. If a dose of bicalutamide tablets is missed, take the next dose at the scheduled time. Do not take the missed dose and do not double the next dose.

#### 2.2 Dosage Adjustment in Renal Impairment

No dosage adjustment is necessary for patients with renal impairment [*see Use in Specific Populations (8.7)*].

#### 2.3 Dosage Adjustment in Hepatic Impairment

No dosage adjustment is necessary for patients with mild to moderate hepatic impairment. In patients with severe liver impairment (n = 4), although there was a 76% increase in the half-life (5.9 and 10.4 days for normal and impaired patients, respectively) of the active enantiomer of bicalutamide no dosage adjustment is necessary [*see Use in Specific Populations (8.6)*].

### 3 DOSAGE FORMS AND STRENGTHS

Bicalutamide tablets 50 mg for oral administration.

### 4 CONTRAINDICATIONS

#### 4.1 Hypersensitivity

Bicalutamide tablets are contraindicated in any patient who has shown a hypersensitivity reaction to the drug or any of the tablet's components. Hypersensitivity reactions including angioneurotic edema and urticaria have been reported [*see Adverse Reactions (6.2)*].

#### 4.2 Women

Bicalutamide tablets have no indication for women, and should not be used in this population.

#### 4.3 Pregnancy

Bicalutamide tablets may cause fetal harm when administered to a pregnant woman. Bicalutamide tablets are contraindicated in women, including those who are or may become pregnant. There are no studies in pregnant women using bicalutamide tablets. If this drug is used during pregnancy, or if the patient becomes pregnant while taking this drug, the patient should be appraised of the potential hazard to the fetus [*see Use in Specific Populations (8.1)*].

5 WARNINGS AND PRECAUTIONS

5.1 Hepatitis

Cases of death or hospitalization due to severe liver injury (hepatic failure) have been reported post-marketing in association with the use of bicalutamide tablets. Hepatotoxicity in these reports generally occurred within the first 3 to 4 months of treatment. Hepatitis or marked increases in liver enzymes leading to drug discontinuation occurred in approximately 1% of bicalutamide tablets patients in controlled clinical trials.

Serum transaminase levels should be measured prior to starting treatment with bicalutamide tablets, at regular intervals for the first 4 months of treatment, and periodically thereafter. If clinical symptoms or signs suggestive of liver dysfunction occur (e.g., nausea, vomiting, abdominal pain, fatigue, anorexia, “flu-like” symptoms, dark urine, jaundice, or right upper quadrant tenderness), the serum transaminases, in particular the serum ALT, should be measured immediately. If at any time a patient has jaundice, or their ALT rises above two times the upper limit of normal, bicalutamide tablets should be immediately discontinued with close follow-up of liver function.

5.2 Gynecomastia and Breast Pain

In clinical trials with bicalutamide tablets 150 mg as a single agent for prostate cancer, gynecomastia and breast pain have been reported in up to 38% and 39% of patients, respectively.

5.3 Glucose Tolerance

A reduction in glucose tolerance has been observed in males receiving LHRH agonists. This may manifest as diabetes or loss of glycemic control in those with pre-existing diabetes. Consideration should therefore be given to monitoring blood glucose in patients receiving bicalutamide tablets in combination with LHRH agonists.

5.4 Laboratory Tests

Regular assessments of serum Prostate Specific Antigen (PSA) may be helpful in monitoring the patient’s response. If PSA levels rise during bicalutamide tablets therapy, the patient should be evaluated for clinical progression. For patients who have objective progression of disease together with an elevated PSA, a treatment-free period of antiandrogen, while continuing the LHRH analog, may be considered.

6 ADVERSE REACTIONS

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

6.1 Clinical Trials Experience

In patients with advanced prostate cancer treated with bicalutamide tablets in combination with an LHRH analog, the most frequent adverse reaction was hot flashes (53%).

In the multicenter, double-blind, controlled clinical trial comparing bicalutamide tablets 50 mg once daily with flutamide 250 mg 3 times a day, each in combination with an LHRH analog, the following adverse reactions with an incidence of 5% or greater, regardless of causality, have been reported.

Table 1. Incidence of Adverse Reactions (≥ 5% in Either Treatment Group) Regardless of Causality

	Treatment Group
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	Number of Patients (%)			
Body System Adverse Reaction	Bicalutamide Plus LHRH Analog (n = 401)		Flutamide Plus LHRH Analog (n = 407)	
Body as a Whole				
Pain (General)	142	(35)	127	(31)
Back Pain	102	(25)	105	(26)
Asthenia	89	(22)	87	(21)
Pelvic Pain	85	(21)	70	(17)
Infection	71	(18)	57	(14)
Abdominal Pain	46	(11)	46	(11)
Chest Pain	34	(8)	34	(8)
Headache	29	(7)	27	(7)
Flu Syndrome	28	(7)	30	(7)
Cardiovascular				
Hot Flashes	211	(53)	217	(53)
Hypertension	34	(8)	29	(7)
Digestive				
Constipation	87	(22)	69	(17)
Nausea	62	(15)	58	(14)
Diarrhea	49	(12)	107	(26)
Increased Liver Enzyme Test	30	(7)	46	(11)
Dyspepsia	30	(7)	23	(6)
Flatulence	26	(6)	22	(5)
Anorexia	25	(6)	29	(7)
Vomiting	24	(6)	32	(8)
Hemic and Lymphatic				
Anemia	45	(11)	53	(13)
Metabolic and Nutritional				
Peripheral Edema	53	(13)	42	(10)
Weight Loss	30	(7)	39	(10)
Hyperglycemia	26	(6)	27	(7)
Alkaline Phosphatase Increased	22	(5)	24	(6)
Weight Gain	22	(5)	18	(4)
Musculoskeletal				
Bone Pain	37	(9)	43	(11)
Myasthenia	27	(7)	19	(5)
Arthritis	21	(5)	29	(7)
Pathological Fracture	17	(4)	32	(8)
Nervous System				
Dizziness	41	(10)	35	(9)
Paresthesia	31	(8)	40	(10)
Insomnia	27	(7)	39	(10)
Anxiety	20	(5)	9	(2)
Depression	16	(4)	33	(8)
Respiratory System				
Dyspnea	51	(13)	32	(8)
Cough Increased	33	(8)	24	(6)

Pharyngitis	32	(8)	23	(6)
Bronchitis	24	(6)	22	(3)
Pneumonia	18	(4)	19	(5)
Rhinitis	15	(4)	22	(5)
<b>Skin and Appendages</b>				
Rash	35	(9)	30	(7)
Sweating	25	(6)	20	(5)
<b>Urogenital</b>				
Nocturia	49	(12)	55	(14)
Hematuria	48	(12)	26	(6)
Urinary Tract Infection	35	(9)	36	(9)
Gynecomastia	36	(9)	30	(7)
Impotence	27	(7)	35	(9)
Breast Pain	23	(6)	15	(4)
Urinary Frequency	23	(6)	29	(7)
Urinary Retention	20	(5)	14	(3)
Urinary Impaired	19	(5)	15	(4)
Urinary Incontinence	15	(4)	32	(8)

Other adverse reactions (greater than or equal to 2%, but less than 5%) reported in the bicalutamide tablets-LHRH analog treatment group are listed below by body system and are in order of decreasing frequency within each body system regardless of causality.

**Body as a Whole:** neoplasm; neck pain; fever; chills; sepsis; hernia; cyst

**Cardiovascular:** angina pectoris; congestive heart failure; myocardial infarct; heart arrest; coronary artery disorder; syncope

**Digestive:** melena; rectal hemorrhage; dry mouth; dysphagia; gastrointestinal disorder; periodontal abscess; gastrointestinal carcinoma

**Metabolic and Nutritional:** edema; BUN increased; creatinine increased; dehydration; gout; hypercholesteremia

**Musculoskeletal:** myalgia; leg cramps

**Nervous:** hypertonia; confusion; somnolence; libido decreased; neuropathy; nervousness

**Respiratory:** lung disorder; asthma; epistaxis; sinusitis

**Skin and Appendages:** dry skin; alopecia; pruritus; herpes zoster; skin carcinoma; skin disorder

**Special Senses:** cataract specified

**Urogenital:** dysuria; urinary urgency; hydronephrosis; urinary tract disorder

**Abnormal Laboratory Test Values:** Laboratory abnormalities including elevated AST, ALT, bilirubin, BUN, and creatinine and decreased hemoglobin and white cell count have been reported in both bicalutamide tablets-LHRH analog treated and flutamide-LHRH analog treated patients.

## 6.2 Postmarketing Experience

The following adverse reactions have been identified during postapproval use of bicalutamide tablets. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

**Respiratory Disorders:** Interstitial lung disease (some fatal) including interstitial pneumonitis and pulmonary fibrosis, most often at doses greater than 50 mg

## **7 DRUG INTERACTIONS**

Clinical studies have not shown any drug interactions between bicalutamide and LHRH analogs (goserelin or leuprolide). There is no evidence that bicalutamide induces hepatic enzymes.

*In vitro* studies have shown that R-bicalutamide is an inhibitor of CYP 3A4 with lesser inhibitory effects on CYP 2C9, 2C19 and 2D6 activity. Clinical studies have shown that with co-administration of bicalutamide tablets, mean midazolam (a CYP 3A4 substrate) levels may be increased 1.5 fold (for  $C_{\max}$ ) and 1.9 fold (for AUC). Hence, caution should be exercised when bicalutamide tablets are co-administered with CYP 3A4 substrates.

*In vitro* protein-binding studies have shown that bicalutamide can displace coumarin anticoagulants from binding sites. Prothrombin times should be closely monitored in patients already receiving coumarin anticoagulants who are started on bicalutamide tablets and adjustment of the anticoagulant dose may be necessary.

## **8 USE IN SPECIFIC POPULATIONS**

### **8.1 Pregnancy**

#### ***Teratogenic Effects. Pregnancy Category X***

*[see Contraindications (4.3)]*

Based on its mechanism of action, bicalutamide tablets may cause fetal harm when administered to a pregnant woman. Bicalutamide tablets are contraindicated in women, including those who are or may become pregnant. If this drug is used during pregnancy, or if the patient becomes pregnant while taking this drug, the patient should be apprised of the potential hazard to a fetus.

While there are no human data on the use of bicalutamide tablets in pregnancy and bicalutamide tablets are not for use in women, it is important to know that maternal use of an androgen receptor inhibitor could affect development of the fetus.

In animal reproduction studies, male offspring of rats receiving doses of 10 mg/kg/day (approximately 2/3 of clinical exposure at the recommended dose) and above, were observed to have reduced anogenital distance and hypospadias. These pharmacological effects have been observed with other antiandrogens. No other teratogenic effects were observed in rabbits receiving doses up to 200 mg/kg/day (approximately 1/3 of clinical exposure at the recommended dose) or rats receiving doses up to 250 mg/kg/day (approximately 2 times the clinical exposure at the recommended dose).

### **8.3 Nursing Mothers**

Bicalutamide tablets are not indicated for use in women.

### **8.4 Pediatric Use**

The safety and effectiveness of bicalutamide in pediatric patients have not been established.

Bicalutamide orodispersible tablet was studied in combination with anastrozole orodispersible tablet in an open-label, non-comparative, multi-center study that assessed the efficacy and safety of this combination regimen over 12 months in the treatment of gonadotropin-independent precocious puberty in boys with familial male-limited precocious puberty, also known as testotoxicosis. Patients were enrolled in the study if they had a baseline age  $\geq 2$  years and a diagnosis of testotoxicosis based on clinical features of progressive precocious puberty, symmetrical testicular enlargement, advanced bone age, pubertal levels of serum testosterone, prepubertal pattern of gonadotropin secretion following a GnRH stimulation test, and absence of other clinical and biochemical causes of testosterone excess.

Thirteen out of the 14 patients enrolled completed 12 months of combination treatment (one patient was lost to follow-up). If central precocious puberty (CPP) developed an LHRH analog was to be added. Four patients were diagnosed with CPP during the 12-month study and received LHRH analog treatment and two additional patients were diagnosed at the end of the 12 months and received treatment subsequently. Mean  $\pm$  SD characteristics at baseline were as follows: chronological age:  $3.9 \pm 1.9$  years; bone age  $8.8 \pm 2.5$ ; bone age/chronological age ratio:  $2.06 \pm 0.51$ ; growth rate (cm/yr):  $10.81 \pm 4.22$ ; growth rate standard deviation score (SDS):  $0.41 \pm 1.36$ .

The starting bicalutamide dose was 12.5 mg. Bicalutamide was titrated in each patient until steady-state R-bicalutamide (the active isomer of bicalutamide) trough plasma concentration reached 5 to 15 mcg/mL, which is the range of therapeutic concentrations achieved in adults with prostate cancer following the administration of the currently approved bicalutamide dose of 50 mg. The starting daily dose of anastrozole was 0.5 mg. Anastrozole was independently titrated in each patient until it reached at steady-state a serum estradiol concentration of  $< 10$  pmol/L (2.7 pg/mL). The following ascending doses were used for bicalutamide: 12.5 mg, 25 mg, 50 mg, and 100 mg. For anastrozole there were two ascending doses: 0.5 mg and 1 mg. At the end of the titration phase one patient was on 12.5 mg bicalutamide, eight patients were on 50 mg bicalutamide, and four patients were on 100 mg bicalutamide; 10 patients were on 0.5 mg anastrozole and three patients were on 1 mg anastrozole. In the majority of patients, steady-state trough concentrations of R-bicalutamide appeared to be attained by Day 21 with once daily dosing. Steady-state trough plasma anastrozole concentrations appeared to be attained by Day 8.

The primary efficacy analysis of the study was to assess the change in growth rate after 12 months of treatment, relative to the growth rate during the  $\geq 6$  months prior to entering the study. Pre-study growth rates were obtained retrospectively. There was no statistical evidence that the growth rate was reduced during treatment. During bicalutamide/anastrozole treatment the mean growth rate (cm/yr) decreased by 1.6 cm/year, 95% CI (- 4.7 to 1.5)  $p = 0.28$ ; the mean growth rate SDS decreased by 0.1 SD, 95% CI (- 1.2 to 1)  $p = 0.88$ . Table 2 shows descriptive data for growth rates for the overall population and for subgroups defined by history of previous treatment for testotoxicosis with ketoconazole, spironolactone, anastrozole or other aromatase inhibitors.

**Table 2. Growth Rates**

Endpoint	Analysis Population	Pre-study Mean	Change from Pre-study to 12 Months			% Patients with Growth Reduction*
			Mean	Median	(Min, Max)	
Growth Rate (cm/yr)	All Treated (n = 13)	10.8	- 1.6	- 2.8	(- 7.4, 8.4)	9/13 (69%)
	PT <sup>†</sup> (n = 6)	10.3	- 0.2	- 2.6 <sup>‡</sup>	(- 7.2, 8.4)	4/6 (67%)
	NPT <sup>§</sup> (n = 7)	11.2	- 2.8	- 2.8	(- 7.4, 1.1)	5/7 (71%)
Growth Rate (SD Units)	All Treated (n = 13)	0.4	- 0.1	- 0.4	(- 2.7, 3.5)	9/13 (69%)
	PT <sup>†</sup> (n = 6)	- 0.1	+ 0.7	- 0.2 <sup>‡</sup>	(- 1.6, 3.5)	4/6 (67%)
	NPT <sup>§</sup> (n = 7)	0.8	- 0.7	- 0.4	(-2.7, 0.5)	5/7 (71%)

\* Change compared to pre-study growth rate

<sup>†</sup> PT = Previous treatment for testotoxicosis with ketoconazole, spironolactone, anastrozole or other aromatase inhibitors

<sup>‡</sup> Median calculated as midpoint of 3<sup>rd</sup> and 4<sup>th</sup> ranked observations



§ NPT = no previous treatment for testotoxicosis with ketoconazole, spironolactone, anastrozole or other aromatase inhibitors

Total testosterone concentrations increased by a mean of 5 mmol/L over the 12 months of treatment from a baseline mean of 10 mmol/L. Estradiol concentrations were at or below the level of quantification (9.81 pmol/L) for 11 of 12 patients after 12 months of treatment. Six of the 12 patients started treatment at an estradiol concentration below the level of quantification.

There were no deaths, serious adverse events, or discontinuations due to adverse events during the study. Of the 14 patients exposed to study treatment, 13 (92.9%) experienced at least one adverse event. The most frequently reported (> 3 patients) adverse events were gynecomastia (7/14, 50%), central precocious puberty (6/14, 43%), vomiting (5/14, 36%), headache (3/14, 21%), pyrexia (3/14, 21%) and upper respiratory tract infection (3/14, 21%). Adverse reactions considered possibly related to bicalutamide by investigators included gynecomastia (6/14, 43%), central precocious puberty (2/14, 14%), breast tenderness (2/14, 14%), breast pain (1/14, 7%), asthenia (1/14, 7%), increased alanine aminotransferase [ALT] (1/14, 7%), increased aspartate aminotransferase [AST] (1/14, 7%), and musculoskeletal chest pain (1/14, 7%). Headache was the only adverse reaction considered possibly related to anastrozole by investigators. For the patient who developed elevated ALT and AST, the elevation was < 3X ULN, and returned to normal without stopping treatment; there was no concomitant elevation in total bilirubin.

## 8.5 Geriatric Use

In two studies in patients given 50 mg or 150 mg daily, no significant relationship between age and steady-state levels of total bicalutamide or the active R-enantiomer has been shown.

## 8.6 Hepatic Impairment

Bicalutamide tablets should be used with caution in patients with moderate-to-severe hepatic impairment. Bicalutamide is extensively metabolized by the liver. Limited data in subjects with severe hepatic impairment suggest that excretion of bicalutamide may be delayed and could lead to further accumulation. Periodic liver function tests should be considered for hepatic-impaired patients on long-term therapy [see *Warnings and Precautions* (5.1)].

No clinically significant difference in the pharmacokinetics of either enantiomer of bicalutamide was noted in patients with mild-to-moderate hepatic disease as compared to healthy controls. However, the half-life of the R-enantiomer was increased approximately 76% (5.9 and 10.4 days for normal and impaired patients, respectively) in patients with severe liver disease (n = 4).

## 8.7 Renal Impairment

Renal impairment (as measured by creatinine clearance) had no significant effect on the elimination of total bicalutamide or the active R-enantiomer.

## 8.8 Women

Bicalutamide has not been studied in women.

## 10 OVERDOSAGE

Long-term clinical trials have been conducted with dosages up to 200 mg of bicalutamide tablets daily and these dosages have been well tolerated. A single dose of bicalutamide tablets that results in symptoms of an overdose considered to be life threatening has not been established.

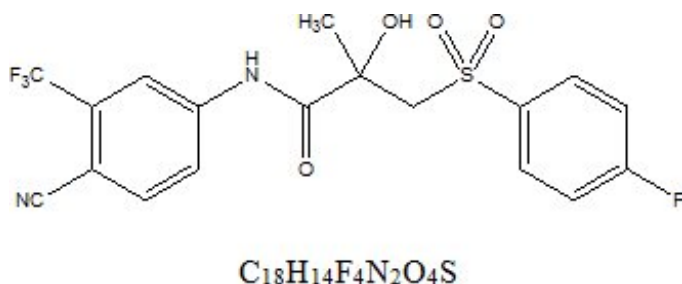
There is no specific antidote; treatment of an overdose should be symptomatic.

In the management of an overdose with bicalutamide tablets, vomiting may be induced if the patient is alert. It should be remembered that, in this patient population, multiple drugs may have been taken.

Dialysis is not likely to be helpful since bicalutamide tablets are highly protein bound and are extensively metabolized. General supportive care, including frequent monitoring of vital signs and close observation of the patient, is indicated.

## 11 DESCRIPTION

Bicalutamide tablets, USP contain 50 mg of bicalutamide, a non-steroidal androgen receptor inhibitor with no other known endocrine activity. The chemical name is propanamide, (+-)-N-[4-cyano-3-(trifluoromethyl)phenyl]-3-[(4-fluorophenyl)sulfonyl]-2-hydroxy-2-methylpropanamide. The structural and molecular formulas are:



Bicalutamide has a molecular weight of 430.37. The pKa' is approximately 12. Bicalutamide, USP is a white crystalline powder which is practically insoluble in water at 37°C (5 mg per 1000 mL), slightly soluble in chloroform and absolute ethanol, sparingly soluble in methanol, and soluble in acetone and tetrahydrofuran.

Bicalutamide tablets are a racemate with its antiandrogenic activity being almost exclusively exhibited by the R-enantiomer of bicalutamide; the S-enantiomer is essentially inactive.

The inactive ingredients of bicalutamide tablets are colloidal silicon dioxide, hypromellose, lactose monohydrate, magnesium stearate, polydextrose, polyethylene glycol, povidone, sodium lauryl sulfate, sodium starch glycolate, titanium dioxide and triacetin.

## 12 CLINICAL PHARMACOLOGY

### 12.1 Mechanism of Action

Bicalutamide tablets are a non-steroidal androgen receptor inhibitor. It competitively inhibits the action of androgens by binding to cytosol androgen receptors in the target tissue. Prostatic carcinoma is known to be androgen sensitive and responds to treatment that counteracts the effect of androgen and/or removes the source of androgen.

When bicalutamide tablets are combined with luteinizing hormone releasing hormone (LHRH) analog therapy, the suppression of serum testosterone induced by the LHRH analog is not affected. However, in clinical trials with bicalutamide tablets as a single agent for prostate cancer, rises in serum testosterone and estradiol have been noted.

In a subset of patients who have been treated with bicalutamide tablets and an LHRH agonist, and who discontinue bicalutamide tablets therapy due to progressive advanced prostate cancer, a reduction in Prostate Specific Antigen (PSA) and/or clinical improvement (antiandrogen withdrawal phenomenon) may be observed.

### 12.3 Pharmacokinetics

#### *Absorption*

Bicalutamide is well-absorbed following oral administration, although the absolute bioavailability is

unknown. Co-administration of bicalutamide with food has no clinically significant effect on rate or extent of absorption.

### ***Distribution***

Bicalutamide is highly protein-bound (96%) [see *Drug Interactions* (7)].

### ***Metabolism/Elimination***

Bicalutamide undergoes stereospecific metabolism. The S (inactive) isomer is metabolized primarily by glucuronidation. The R (active) isomer also undergoes glucuronidation but is predominantly oxidized to an inactive metabolite followed by glucuronidation. Both the parent and metabolite glucuronides are eliminated in the urine and feces. The S-enantiomer is rapidly cleared relative to the R-enantiomer, with the R-enantiomer accounting for about 99% of total steady-state plasma levels.

Pharmacokinetics of the active enantiomer of bicalutamide tablets in normal males and patients with prostate cancer are presented in Table 3.

**Table 3**

<b>Parameter</b>	<b>Mean</b>	<b>Standard Deviation</b>
Normal Males (n = 30)		
Apparent Oral Clearance (L/hr)	0.320	0.103
Single Dose Peak Concentration (mcg/mL)	0.768	0.178
Single Dose Time to Peak Concentration (hours)	31.3	14.6
Half-life (days)	5.8	2.29
Patients with Prostate Cancer (n = 40)		
C <sub>ss</sub> (mcg /mL)	8.939	3.504

## **13 NONCLINICAL TOXICOLOGY**

### **13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility**

Two-year oral carcinogenicity studies were conducted in both male and female rats and mice at doses of 5, 15 or 75 mg/kg/day of bicalutamide. A variety of tumor target organ effects were identified and were attributed to the antiandrogenicity of bicalutamide, namely, testicular benign interstitial (Leydig) cell tumors in male rats at all dose levels (the steady-state plasma concentration with the 5 mg/kg/day dose is approximately 2/3 human therapeutic concentrations\*) and uterine adenocarcinoma in female rats at 75 mg/kg/day (approximately 1 1/2 times the human therapeutic concentrations). There is no evidence of Leydig cell hyperplasia in patients; uterine tumors are not relevant to the indicated patient population.

A small increase in the incidence of hepatocellular carcinoma in male mice given 75 mg/kg/day of bicalutamide (approximately 4 times human therapeutic concentrations\*) and an increased incidence of benign thyroid follicular cell adenomas in rats given 5 mg/kg/day (approximately 2/3 human therapeutic concentrations\*) and above were recorded. These neoplastic changes were progressions of non-neoplastic changes related to hepatic enzyme induction observed in animal toxicity studies. Enzyme induction has not been observed following bicalutamide administration in man. There were no tumorigenic effects suggestive of genotoxic carcinogenesis.

A comprehensive battery of both *in vitro* and *in vivo* genotoxicity tests (yeast gene conversion, Ames, E. coli, CHO/HGPRT, human lymphocyte cytogenetic, mouse micronucleus, and rat bone marrow cytogenetic tests) has demonstrated that bicalutamide tablets do not have genotoxic activity.

Administration of bicalutamide tablets may lead to inhibition of spermatogenesis. The long-term effects of bicalutamide tablets on male fertility have not been studied.

In male rats dosed at 250 mg/kg/day (approximately 2 times human therapeutic concentrations\*), the precoital interval and time to successful mating were increased in the first pairing but no effects on fertility following successful mating were seen. These effects were reversed by 7 weeks after the end of an 11-week period of dosing.

No effects on female rats dosed at 10, 50 and 250 mg/kg/day (approximately 2/3, 1 and 2 times human therapeutic concentrations, respectively\*) or their female offspring were observed. Administration of bicalutamide to pregnant females resulted in feminization of the male offspring leading to hypospadias at all dose levels. Affected male offspring were also impotent.

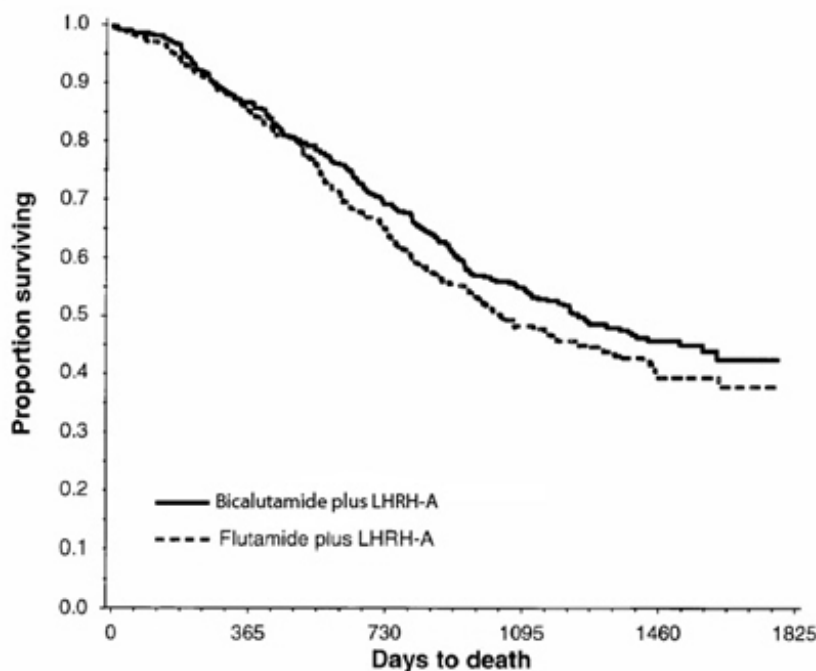
\*Based on a maximum dose of 50 mg/day of bicalutamide for an average 70 kg patient.

## 14 CLINICAL STUDIES

### 14.1 Bicalutamide Tablets 50 mg Daily in Combination with an LHRH-A

In a multicenter, double-blind, controlled clinical trial, 813 patients with previously untreated advanced prostate cancer were randomized to receive bicalutamide tablets 50 mg once daily (404 patients) or flutamide 250 mg (409 patients) 3 times a day, each in combination with LHRH analogs (either goserelin acetate implant or leuprolide acetate depot).

In an analysis conducted after a median follow-up of 160 weeks was reached, 213 (52.7%) patients treated with bicalutamide tablets-LHRH analog therapy and 235 (57.5%) patients treated with flutamide-LHRH analog therapy had died. There was no significant difference in survival between treatment groups (see Figure 1). The hazard ratio for time to death (survival) was 0.87 (95% confidence interval 0.72 to 1.05).



**Figure 1 - The Kaplan-Meier Probability of Death for Both Antiandrogen Treatment Groups.**

Figure 1 - The Kaplan-Meier Probability of Death for Both Antiandrogen Treatment Groups.

There was no significant difference in time to objective tumor progression between treatment groups (see Figure 2). Objective tumor progression was defined as the appearance of any bone metastases or the worsening of any existing bone metastases on bone scan attributable to metastatic disease, or an increase by 25% or more of any existing measurable extraskelatal metastases. The hazard ratio for time to progression of bicalutamide tablets plus LHRH analog to that of flutamide plus LHRH analog was 0.93 (95% confidence interval, 0.79 to 1.10).

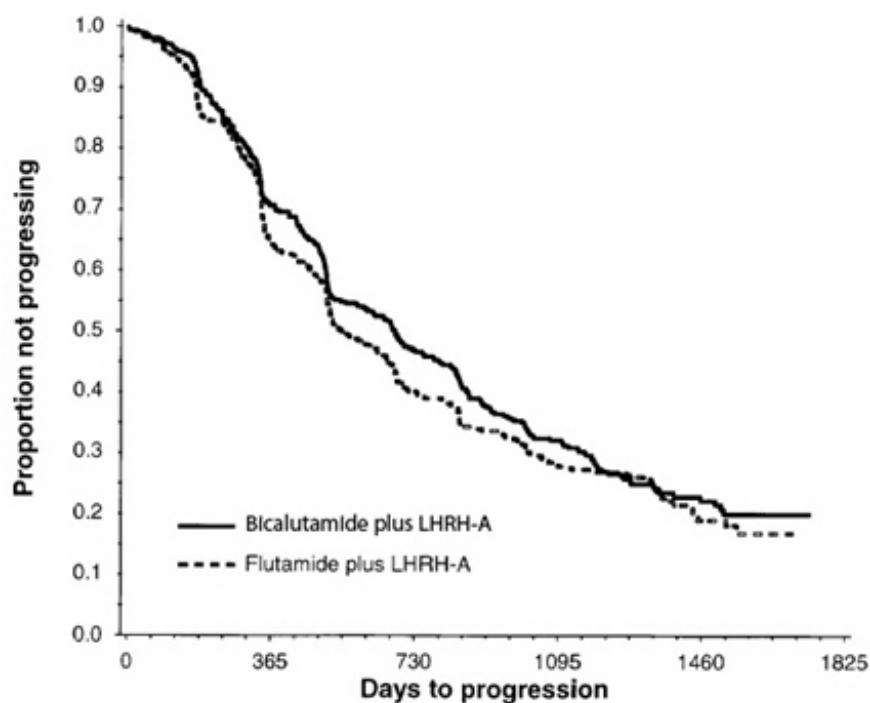


Figure 2 - Kaplan-Meier Curve for Time to Progression for Both Antiandrogen Treatment Groups.

Figure 2 - Kaplan-Meier Curve for Time to Progression for Both Antiandrogen Treatment Groups.

Quality of life was assessed with self-administered patient questionnaires on pain, social functioning, emotional well being, vitality, activity limitation, bed disability, overall health, physical capacity, general symptoms, and treatment related symptoms. Assessment of the Quality of Life questionnaires did not indicate consistent significant differences between the two treatment groups.

#### 14.2 Safety Data from Clinical Studies Using Bicalutamide Tablets 150 mg

**Bicalutamide tablets 150 mg are not approved for use either alone or with other treatments.**

Two identical multicenter, randomized, open-label trials comparing bicalutamide tablets 150 mg daily monotherapy to castration were conducted in patients that had locally advanced (T3-4, NX, MO) or metastatic (M1) prostate cancer.

##### ***Monotherapy***

##### ***M1 Group***

Bicalutamide tablets 150 mg daily are not approved for use in patients with M1 cancer of the prostate. Based on an interim analysis of the two trials for survival, the Data Safety Monitoring Board recommended that bicalutamide tablets treatment be discontinued in the M1 patients because the risk of death was 25% (HR 1.25, 95% CI 0.87 to 1.81) and 31% (HR 1.31, 95% CI 0.97 to 1.77) higher in the

bicalutamide tablets treated group compared to that in the castrated group, respectively.

#### *Locally Advanced (T3-4, NX, MO) Group*

Bicalutamide tablets 150 mg daily are not approved for use in patients with locally advanced (T3-4, NX, MO) cancer of the prostate. Following discontinuation of all M1 patients, the trials continued with the T3-4, NX, MO patients until study completion. In the larger trial (N = 352), the risk of death was 25% (HR 1.25, 95% CI 0.92 to 1.71) higher in the bicalutamide tablets group and in the smaller trial (N = 140), the risk of death was 36% (HR 0.64, 95% CI, 0.39 to 1.03) lower in the bicalutamide tablets group.

In addition to the above two studies, there are three other on-going clinical studies that provide additional safety information for bicalutamide tablets 150 mg, a dose that is not approved for use. These are three multicenter, randomized, double-blind, parallel group trials comparing bicalutamide tablets 150 mg daily monotherapy (adjuvant to previous therapy or under watchful waiting) with placebo, for death or time to disease progression, in a population of 8,113 patients with localized or locally advanced prostate cancer.

Bicalutamide tablets 150 mg daily are not approved for use as therapy for patients with localized prostate cancer who are candidates for watchful waiting. Data from a planned subgroup analysis of two of these trials in 1,627 patients with localized prostate cancer who were under watchful waiting, revealed a trend toward decreased survival in the bicalutamide tablets arm after a median follow-up of 7.4 years. There were 294 (37.7%) deaths in the bicalutamide tablets treated patients versus 279 (32.9%) deaths in the placebo treated patients (localized watchful waiting group) for a hazard ratio of 1.16 (95% CI 0.99 to 1.37).

## **16 HOW SUPPLIED/STORAGE AND HANDLING**

Bicalutamide Tablets, USP are available containing 50 mg of bicalutamide, USP.

The 50 mg tablets are white film-coated, round, unscored tablets debossed with **M** on one side of the tablet and **C17** on the other side. They are available as follows:

NDC 0378-7017-93  
bottles of 30 tablets

### **16.1 Storage and Handling**

**Store at 20° to 25°C (68° to 77°F). [See USP Controlled Room Temperature.]**

Dispense in a tight, light-resistant container as defined in the USP using a child-resistant closure.

**PHARMACIST:** Dispense a Patient Information Leaflet with each prescription.

## **17 PATIENT COUNSELING INFORMATION**

Advise the patient to read the FDA-approved patient labeling (Patient Information).

Patients should be informed that therapy with bicalutamide tablets and the LHRH analog should be started at the same time and that they should not interrupt or stop taking these medications without consulting their physician.

During treatment with bicalutamide tablets, somnolence has been reported, and those patients who experience this symptom should observe caution when driving or operating machines.

Patients should be informed that diabetes, or loss of glycemic control in patients with pre-existing diabetes has been reported during treatment with LHRH agonists. Consideration should therefore be given to monitoring blood glucose in patients receiving bicalutamide tablets in combination with LHRH agonists.

Patients should be informed that cases of photosensitivity have been reported during treatment with bicalutamide tablets and that they should avoid direct exposure to excessive sunlight or UV-light exposure. Consideration should be given to the use of sunscreen.

## **PATIENT INFORMATION**

### **BICALUTAMIDE TABLETS, USP**

**(bye'' ka loo' ta mide)**

**50 mg**

Read this Patient Information before you start taking bicalutamide tablets and each time you get a refill. There may be new information. This leaflet does not take the place of talking with your healthcare provider about your medical condition or your treatment.

#### **What are bicalutamide tablets?**

Bicalutamide tablets are a prescription medicine called an androgen receptor inhibitor, used in combination with lutenizing hormone-releasing hormone (LHRH) medicines to treat stage D<sub>2</sub> metastatic prostate cancer.

Bicalutamide tablets are not for use in women.

It is not known if bicalutamide tablets are safe and effective in children.

#### **Who should not take bicalutamide tablets?**

Do not take bicalutamide tablets if you:

- are a woman.
- are pregnant or may become pregnant. Bicalutamide may harm your unborn baby
- are allergic to any of the ingredients in bicalutamide tablets. See the end of this leaflet for a complete list of ingredients in bicalutamide tablets. Get medical help right away if you get any of the following symptoms of an allergic reaction: itching, hives (raised bumps), swelling of the face, lips or tongue, trouble breathing or swallowing.

#### **What should I tell my healthcare provider before taking bicalutamide tablets?**

**Before you take bicalutamide tablets, tell your healthcare provider about all your medical conditions, including if you:**

- have liver problems.
- take a medicine to thin your blood. Ask your healthcare provider or pharmacist if you are not sure if your medicine is a blood thinner.
- have diabetes (poor blood sugar control has been reported in people taking bicalutamide tablets in combination with LHRH medicines).

**Tell your healthcare provider about all the medicines you take**, including prescription and over the counter medicines, vitamins and herbal supplements. Bicalutamide tablets and other medicines may affect each other causing side effects. Bicalutamide tablets may affect the way other medicines work, and other medicines may affect how bicalutamide tablets work.

Know the medicines you take. Keep a list of your medicines with you to show your healthcare providers when you get a new medicine.

#### **How should I take bicalutamide tablets?**

- Take bicalutamide tablets exactly as your healthcare provider tells you to take it.
- Take bicalutamide tablets at the same time everyday.

- Your treatment with bicalutamide tablets should start at the same time as your treatment with the LHRH medicine.
- If you miss a dose do not take an extra dose, take the next dose at your regular time. Do not take two doses at the same time.
- Bicalutamide tablets can be taken with or without food.
- If you take too many bicalutamide tablets, call your healthcare provider or Poison Control Center or go to the nearest hospital emergency room right away.
- Do not stop taking bicalutamide tablets unless your healthcare provider tells you.
- Your healthcare provider may do blood tests while you take bicalutamide tablets.
- Your prostate cancer may get worse while taking bicalutamide tablets in combination with LHRH medicines. Regular monitoring of your prostate cancer with your healthcare provider is important to determine if your disease is worse.

### **What should I avoid while taking bicalutamide tablets?**

Do not drive, operate machinery, or do other dangerous activities until you know how bicalutamide tablets affect you.

Some people have had skin sensitivity to sunlight while taking bicalutamide tablets. You should avoid sunlight or sunlamps and tanning beds and consider using sunscreen while being treated with bicalutamide tablets.

### **What are the possible side effects of bicalutamide tablets?**

#### **Bicalutamide tablets may cause serious side effects, including:**

- Liver problems, including liver failure that may need to be treated in a hospital or that may lead to death. Your healthcare provider should do blood tests to check your liver function before and during treatment with bicalutamide tablets. Tell your healthcare provider right away if you get any of these symptoms of a liver problem during treatment:
  - yellowing of the skin and eyes (jaundice)
  - dark urine
  - right upper stomach pain
  - nausea
  - vomiting
  - tiredness
  - loss of appetite
  - chills
  - fever
- Trouble breathing with or without a cough or fever. Some people taking bicalutamide tablets get an inflammation in the lungs called interstitial lung disease.
- An allergic reaction. Symptoms of an allergic reaction include: itching of the skin, hives (raised bumps), swelling of the face, lips, tongue, throat or trouble swallowing.
- Enlargement of breast (gynecomastia) and breast pain.
- Poor blood sugar control can happen in people who take bicalutamide tablets in combination with LHRH medicines. Your healthcare provider may check your blood sugar during bicalutamide tablets therapy.

The most common side effects of bicalutamide tablets include:



- o hot flashes, or short periods of feeling warm and sweating
- o whole body pain in your back, pelvis, stomach
- o feeling weak
- o constipation
- o waking from sleep to urinate at night
- o a decrease in red blood cells (anemia)
- o infection
- o nausea
- o swelling in your ankles, legs or feet
- o diarrhea
- o blood in your urine
- o feeling dizzy

Tell your healthcare provider if you have any side effect that bothers you or that does not go away.

These are not all the possible side effects of bicalutamide tablets. For more information, ask your healthcare provider or pharmacist.

**Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.**

#### **How should I store bicalutamide tablets?**

Store bicalutamide tablets at 20° to 25°C (68° to 77°F).

**Keep bicalutamide tablets and all medicines out of the reach of children.**

#### **General information about the safe and effective use of bicalutamide tablets.**

Medicines are sometimes prescribed for purposes other than those listed in a patient information leaflet. Do not use bicalutamide tablets for a condition for which it was not prescribed. Do not give bicalutamide tablets to other people, even if they have the same symptoms that you have. It may harm them.

This Patient Information leaflet summarizes the most important information about bicalutamide tablets. If you would like more information about bicalutamide tablets talk with your healthcare provider. You can ask your healthcare provider or pharmacist for information about bicalutamide tablets that is written for health professionals. For more information, call Mylan Pharmaceuticals Inc. at 1-877-446-3679 (1-877-4-INFO-RX).

#### **What are the ingredients in bicalutamide tablets, USP?**

**Active ingredients:** bicalutamide, USP

**Inactive ingredients:** colloidal silicon dioxide, hypromellose, lactose monohydrate, magnesium stearate, polydextrose, polyethylene glycol, povidone, sodium lauryl sulfate, sodium starch glycolate, titanium dioxide and triacetin.

This Patient Information has been approved by the U.S. Food and Drug Administration.

**Mylan Pharmaceuticals Inc.**

Morgantown, WV 26505 U.S.A.

REVISED APRIL 2015

BCLT:R4p

**PRINCIPAL DISPLAY PANEL - 50 mg**

**NDC 0378-7017-93**

**Bicalutamide  
Tablets, USP  
50 mg**

**PHARMACIST: Dispense the accompanying  
Patient Information Leaflet to each patient.**

**Rx only    30 Tablets**

Each film-coated tablet contains:  
Bicalutamide, USP        50 mg

Dispense in a tight, light-resistant  
container as defined in the USP  
using a child-resistant closure.

Keep container tightly closed.

**Keep this and all medication  
out of the reach of children.**

**Store at 20° to 25°C (68° to 77°F).  
[See USP Controlled Room  
Temperature.]**

**Usual Dosage:** See accompanying  
prescribing information.

**Mylan Pharmaceuticals Inc.**  
Morgantown, WV 26505 U.S.A.

**Mylan.com**

**RM7017H3**

3  
0378-7017-93  
4

Each film-coated tablet contains:  
Bicalutamide, USP        50 mg

**NDC 0378-7017-93**

**Bicalutamide**  
Tablets, USP

**50 mg**

PHARMACIST: Dispense the accompanying  
Patient Information Leaflet to each patient.

**Mylan®**

**Rx only    30 Tablets**

Dispense in a tight, light-resistant  
container as defined in the USP  
using a child-resistant closure.  
Keep container tightly closed.  
**Keep this and all medication  
out of the reach of children.**  
Store at 20° to 25°C (68° to 77°F).  
[See USP Controlled Room  
Temperature.]  
Usual Dosage: See accompanying  
prescribing information.  
Mylan Pharmaceuticals Inc.  
Morgantown, WV 26505 U.S.A.

**Mylan®**  
Mylan.com

RM7017H3

**BICALUTAMIDE**

bicalutamide tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:0378-7017
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Route of Administration		ORAL		DEA Schedule	
Active Ingredient/Active Moiety					
Ingredient Name			Basis of Strength		Strength
Bicalutamide (UNII: A0Z3NAU9DP) (Bicalutamide - UNII:A0Z3NAU9DP)			Bicalutamide		50 mg
Inactive Ingredients					
Ingredient Name					Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)					
HYPROMELLOSES (UNII: 3NXW29V3WO)					
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)					
MAGNESIUM STEARATE (UNII: 70097M6I30)					
POLYDEXTROSE (UNII: VH2XOU12IE)					
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)					
POVIDONES (UNII: FZ989GH94E)					
SODIUM LAURYL SULFATE (UNII: 368GB5141J)					
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)					
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)					
TRIACETIN (UNII: XHX3C3X673)					
Product Characteristics					
Color	WHITE	Score	no score		
Shape	ROUND	Size	6mm		
Flavor		Imprint Code	M;C17		
Contains					
Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:0378-7017-93	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
Marketing Information					
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date	
ANDA	ANDA079185		07/06/2009		
Labeler - Mylan Pharmaceuticals Inc (059295980)					